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USE OF ADENOSINE DEAMINASE INHIBITORS TO TREAT  
SYSTEMIC INFLAMMATORY RESPONSE SYNDROME

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5 This application claims priority to co-pending PCT/US00/13987, filed May 22,  
2000 --and U.S. Serial No. 09/317,678 filed May 24, 1999, now U.S. Patent No.  
6,103,702.--

This invention relates to a new use of adenosine deaminase inhibitors in the  
prevention or treatment of adverse consequences of systemic inflammatory responses  
10 (SIRS). These conditions are ameliorated by increasing the local concentration of  
adenosine in affected regions.

**BACKGROUND OF THE INVENTION**

Conditions resulting in or from a systemic inflammatory response syndrome  
(SIRS) are associated with an exaggerated immune response, oxygen free-radical-  
15 mediated injury, and tissue perfusion maldistribution. Such conditions include endotoxin  
shock, septic shock, sepsis, endotoxemia, septicemia, peritonitis, and adult respiratory  
distress syndrome (ARDS). Current treatment is unsatisfactory. Therapeutic attempts  
to modify cytokine responses during SIRS-related conditions have focussed on antibodies  
to the cytokines or cytokine receptor antagonists. These approaches have proven  
20 unsuccessful because some level of cytokine response is required for survival from SIRS-  
related conditions.

Adenosine has been reported to be an endogenous modulator of inflammation by  
virtue of its effects on stimulated granulocyte function (Cronstein *et al.*, 1986) and on  
macrophage, lymphocyte and platelet function. Adenosine receptor agonists have been  
25 reported to be beneficial in an experimental model of inflammation (Schrier *et al.*, 1990).  
Adenosine and a related analog have been reported to inhibit in vitro production of the  
cytokine, tumor necrosis factor  $\alpha$  (Parmely *et al.*, 1991). Antibodies to TNF- $\alpha$  have not  
been shown to alter mortality in sepsis (Abraham *et al.*, 1998; Cohen *et al.*, 1996; and  
Amiot *et al.*, 1997).

30 Adenosine is an endogenous, ubiquitous molecule that modulates immune  
function, can suppress or increase free-radical production, and produces vasodilation in  
regions wherein adenosine is produced in significant quantities.